

IN VITRO CULTURE OF AQUATIC PLANT
Aglonema simplex

WILLYAH ST. MOHD RAHMI

JABATAN SAINS BIOLOGI
FAKULTI SAINS DAN TEKNOLOGI
KOLEJ UNIVERSITI SAINS DAN TEKNOLOGI MALAYSIA
2004

1100030779

PERPUSTAKAAN KOLEJ UNIVERSITI SAINS & TEKNOLOGI MALAYSIA (KUSTEM)			
Pengarang	Judul	No. Panggilan	
Tarikh	Waktu Pemulangan	Nombor Ahli	Tanda tangan
Wardah bt Mond Pandiar	In vitro culture of aquatic plant	LP 26 PCT 16 5004	
16/8/05	3.00 pm	UK10523	2.
24/7/06	4.00 pm	UK10532	m.
8/8/06	1.20 pts.	UK10504	x
12/8/06	2.00 pm	UK10532	le.
13/8/06	7.00 pm	UK10532	
14/8/06	2.00 pm	UK10532	
	+ 00 hrs		

1100030779

LP 26 FST 3 2004



1100030779

In vitro culture of aquatric plant (*Aglaonema simplex*) / Wardah Mohd. Pauzi.



PERPUSTAKAAN

**KOLEJ UNIVERSITI SAINS & TEKNOLOGI MALAYSIA
21030 KUALA TERENGGANU**

21030 KELAH TERENGGANG
1100030779

Lihat sebelah

HAK MILIK
PERPUSTAKAAN KUSTEM

IN VITRO CULTURE OF AQUATIC PLANT
Aglaonema simplex

By

Wardah bt. Mohd Paudzi

**Research Report submitted in partial fulfilment of
the requirements for the degree of
Bachelor of Science (Biological Sciences)**

**Department of Biological Sciences
Faculty of Science and Technology
KOLEJ UNIVERSITI SAINS DAN TEKNOLOGI MALAYSIA
2004**



JABATAN SAINS BIOLOGI
FAKULTI SAINS DAN TEKNOLOGI
KOLEJ UNIVERSITI SAINS DAN TEKNOLOGI MALAYSIA

PENGAKUAN DAN PENGESAHAN LAPORAN
PROJEK PENYELIDIKAN I DAN II

Adalah ini diakui dan disahkan bahawa laporan penyelidikan bertajuk:

In vitro culture of aquatic plant Aglaonema simplex

oleh Wardah Binti Mohd Paudzi, No. Matrik UK 5642

telah diperiksa dan semua pembetulan yang disarankan telah dilakukan. Laporan ini dikemukakan kepada Jabatan Sains Biologi sebagai memenuhi sebahagian daripada keperluan memperolehi Ijazah Sarjana Muda Sains – Sains Biologi,

Fakulti Sains dan Teknologi, Kolej Universiti Sains dan Teknologi Malaysia.

Disahkan oleh,

Penyelia Utama

Nama: DR. AZIZ BIN AHMAD (Ph.D)
LECTURER
Cop Rasmi: Dept of Biological Sciences
Fakulty of Science and Technology
University Collage of Science
and Technology Malaysia
21030 Kuala Terengganu.

Tarikh: 30/3/2004

.....
Penyelia Kedua (jika ada)

Nama:

Cop Rasmi:

Tarikh:

.....
Ketua Jabatan Sains Biologi

PROF. DR. CHAN ENG HENG
Ketua
Jabatan Sains Biologi
Fakulti Sains dan Teknologi
Kolej Universiti Sains dan Teknologi Malaysia
(KUSTEM)
21030 Kuala Terengganu.

Tarikh: 30/3/2004

Nama:

Cop Rasmi:

ACKNOWLEDGEMENT

First of all, Alhamdulillah, thank God for giving me strength to complete all my work on time. I would like to thank Dr. Aziz Bin Ahmad for helping me doing my project and also being such a great supervisor to me. Thank you also to my parents and to all my family for all the support and encouragement.

Thank also due to Abas, Kak Rokiah, Abang Syed, Chien, Chals, Nisha, Sya and Nyamuk for helping me at Biotech Lab. Last but not least, to my special friend, Yusz, thanks for all your patience and support.

TABLE OF CONTENTS

	Page
ACKNOWLEDGEMENT	ii
LIST OF TABLES	v
LIST OF FIGURES	vi
LIST OF PLATES	vii
LIST OF APPENDICES	viii
LIST OF SYMBOLS	ix
ABSTRACT	x
ABSTRAK	xi
1.0 INTRODUCTION	1
1.1 Objective	3
2.0 LITERATURE REVIEW	
2.1 The <i>Aglaonema simplex</i>	4
2.2 <i>Aglaonema</i> Diseases	5
2.3 Aquarium Plants Industry	6
2.4 Application of Tissue Culture on Aquatic Plant	7
2.5 Culture Medium	8
2.6 The Concept of Plant Growth Regulators	8
2.6.1 Cytokinins	9
3.0 MATERIALS AND METHODS	
3.1 Sources of Explants	11

3.2	Sterilization of Explants	11
3.3	Medium for Culture Establishment	13
3.4	Effect of Medium and Cytokinin On Growth of Cultures	15
3.5	Effect of Cutting Technique on Proliferation of Shoot	16
4.0	RESULTS AND DISCUSSIONS	
4.1	Establishment of Sterile Culture	18
4.2	Effect of Medium and Cytokinin On Growth of Cultures	20
4.3	Effect of Cutting Technique on Proliferation of Shoot	27
5.0	CONCLUSION	32
6.0	LIST OF REFERENCE	33
	APPENDICES	35
	CURRICULUM VITAE	39

LIST OF TABLES

Table	Title	Page
1	Plants have been propagated using tissue culture technique.	7
2	The percentage of Clorox with different times of immersion.	13
3	Concentration of different types of cytokinin.	15
4	The percentage of explants free of contamination after treatment with various concentrations of Clorox at different times of immersion.	18
5	Number of shoot produced from different kind of cutting technique of explants in MS medium with BAP treatment.	27

LIST OF FIGURES

Figure	Title	Page
1	Flow chart showing the experiment 3.4.	16
2	Flow chart showing the experiment 3.5.	17
3	Growth of <i>A. simplex</i> in different types of media with various concentrations of BAP, (a) MS + BAP, (b) B5 + BAP, (c) AS + BAP.	21
4	Growth of <i>A. simplex</i> in different types of media with various concentrations of Kinetin, (a) MS + Kinetin, (b) B5 + Kinetin, (c) AS + Kinetin.	22
5	Growth of <i>A. simplex</i> in different types of media with various concentrations of 2iP, (a) MS + 2iP, (b) B5 + 2iP, (c) AS + 2iP.	24
6	Fresh weight of <i>A. simplex</i> at 28 days with, (a) BAP, (b) Kinetin, (c) 2iP.	25

LIST OF PLATES

Plate	Title	Page
1	(a) <i>A. simplex</i> at Lata Tembakah (b) <i>A. simplex</i> in an aquarium.	12
2	Shoot of <i>A. simplex</i> cultured in MS medium containing 1.0 mg/L of BAP.	14
3	(a) Plant cultured on MS medium supplemented with 5.0 mg/L BAP at 0 days. (b) After 28 days.	26
4	(a) Plantlet cultured on MS medium supplemented with 5.0 mg/L BAP at 0 days. (b) After 28 days.	29
5	(a) Two parts of shoot from longitudinal section used to establish cultures. After 28 days cultured on MS media supplemented with (b) 0.0 mg/L BAP (c) 1.0 mg/L BAP. (d) 3.0 mg/L BAP. (e) 5.0 mg/L BAP.	30
6	(a) Three parts of shoot from cross section used to establish cultures. After 28 days cultured on MS media supplemented with (b) 0.0 mg/L BAP (c) 1.0 mg/L BAP. (d) 3.0 mg/L BAP. (e) 5.0 mg/L BAP.	31

LIST OF APPENDICES

Appendix		Page
1	AS medium.	35
2	Table 6: Fresh weight of <i>A. simplex</i> cultured in MS medium with (a) BAP, (b) Kinetin, (c) 2iP	36
3	Table 7: Fresh weight of <i>A. simplex</i> cultured in B5 medium with (a) BAP, (b) Kinetin, (c) 2iP	37
4	Table 8: Fresh weight of <i>A. simplex</i> cultured in AS medium with (a) BAP, (b) Kinetin, (c) 2iP	38

LIST OF SYMBOLS

mg/L	-	Milligram per liter
BAP	-	Benzylaminopurine
Kin	-	Kinetin
2iP	-	2-Isopentyladenine
HCl	-	Hydrochloric acid
NaOH	-	Natrium hydrochloride
M	-	Molar
v/w	-	Volume per weight
v/v	-	Volume per volume

ABSTRACT

An aquatic plant, *Aglaonema simplex* has been successfully cultured in – vitro using shoot tip in MS medium containing 1.0 mg/L of BAP. The optimum sterilization condition was by using 100% (v/v) of Clorox with 30 minutes of immersion time. The effect of media (MS medium, B5 medium and AS medium) and cytokinin (BAP, Kinetin and 2iP) were examined. The best growth was obtained in MS medium containing 5.0 mg/L of BAP. Three types of explants, which were complete plantlet, shoot in longitudinal section and shoot in cross section were used for induction of shoot proliferation. The highest number of shoot proliferation was obtained using longitudinal section, which cultured in MS medium containing 5.0 mg/L of BAP.

ABSTRAK

Kultur in – vitro tumbuhan akuatik, *Aglaonema simplex* telah berjaya dihasilkan dengan menggunakan pucuk dan dikultur di dalam media MS yang mengandungi 1.0 mg/L BAP. Kadar pensterilan yang optimum didapati dengan menggunakan 100% (v/v) Klorox, direndam selama 30 minit. Kesan media (MS, B5 dan AS) dan sitokinin (BAP, Kinetin dan 2iP) telah dikaji. Pertumbuhan pokok yang terbaik didapati di dalam media MS yang mengandungi 5.0 mg/L BAP. Tiga jenis eksplan iaitu pokok yang sempurna, pucuk pada keratan memanjang dan pucuk pada keratan melintang digunakan untuk mengaruh pertumbuhan pucuk yang baru. Bilangan pucuk yang tertinggi didapati dengan menggunakan pucuk pada keratan memanjang yang dikultur di dalam media MS yang mengandungi 5.0 mg/L BAP.